Welcome to STN International! Enter x:x

LOGINID: SSSPTAU188MXM

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

NEWS 1 Web Page URLs for STN Seminar Schedule - N. America

NEWS 2 "Ask CAS" for self-help around the clock

NEWS 3 JUL 20 Powerful new interactive analysis and visualization software, STN AnaVist, now available

NEWS 4 AUG 11 STN AnaVist workshops to be held in North America

NEWS 5 AUG 30 CA/Caplus -Increased access to 19th century research documents

NEWS 6 AUG 30 CASREACT - Enhanced with displayable reaction conditions

NEWS 7 SEP 09 ACD predicted properties enhanced in REGISTRY/ZREGISTRY

NEWS 8 OCT 03 MATHDI removed from STN

NEWS 9 OCT 04 CA/Caplus-Canadian Intellectual Property Office (CIPO) added to core patent offices

NEWS 10 OCT 06 STN AnaVist workshops to be held in North America

NEWS 11 OCT 13 New CAS Information Use Policies Effective October 17, 2005

NEWS EXPRESS JUNE 13 CURRENT WINDOWS VERSION IS V8.0, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 13 JUNE 2005

NEWS HOURS STN Operating Hours Plus Help Desk Availability

NEWS INTER General Internet Information

NEWS LOGIN Welcome Banner and News Items

NEWS PHONE Direct Dial and Telecommunication Network Access to STN

NEWS WWW CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that specific topic.

All use of STN is subject to the provisions of the STN Customer agreement. Please note that this agreement limits use to scientific research. Use for software development or design or implementation of commercial gateways or other similar uses is prohibited and may result in loss of user privileges and other penalties.

FILE 'HOME' ENTERED AT 18:34:18 ON 14 OCT 2005

=> file ca, biosis,medline
COST IN U.S. DOLLARS

FULL ESTIMATED COST

ST IN U.S. DOLLARS SINCE FILE ENTRY

ENTRY SESSION 0.84 0.84

TOTAL

FILE 'CA' ENTERED AT 18:36:52 ON 14 OCT 2005 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

COPYRIGHT (C) 2005 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'BIOSIS' ENTERED AT 18:36:52 ON 14 OCT 2005 Copyright (c) 2005 The Thomson Corporation

FILE 'MEDLINE' ENTERED AT 18:36:52 ON 14 OCT 2005

```
=> s (betaine hcl) or (betaine hydrochloric acid)
L1
          242 (BETAINE HCL) OR (BETAINE HYDROCHLORIC ACID)
=> s pepsin
        41097 PEPSIN
\Rightarrow s 11 and 12
           13 L1 AND L2
=> d 1-13 ab,bib
    ANSWER 1 OF 13 CA COPYRIGHT 2005 ACS on STN
L3
AB
     The present invention provides a stabilized protonic formulation comprised
     primarily of proteins, enzymes and pH adjusters, all in specific ratios to
     one another, a liquid medium which, when combined to the protonic
     formulation, initiates activation of the amino acids within the protonic
     formulation, and a stabilizing component which stabilizes the amino acids
     during the process of their activation. The optimum ratio of enzyme
     activator formulation to protein mixture is about 1:25, though 1 part enzyme
     activator formulation to 10 to 30 parts protein mixture will function
     suitably for the intended purpose. The enzyme activator formulation is
     optimally comprised of: betaine-HCl 4.0%,
     pepsin 1.5%, trypsin 0.4%, chymotrypsin 0.3%, protease 0.4%,
    bromelain 0.5%, papaya 0.6%, vitamin C 5.0%, lemon powder 0.6%, glutamic
     acid 0.2%, and glycine 86.4%. The protein sources and mixture that works
    best with the enzyme activator formulation includes: whey protein isolate
     30.0%, instant whey concentrate 15.0%, soy protein isolate 25.0%, pea protein
     5.0%, rice protein 5.0%, maltodextrin 15.7%, steviocide 0.3%, french
     vanilla flavor 1.75%, peach mango flavor 0.5%, xanthan gum 0.5%, lecithin
     0.5%, and tricalcium phosphate 0.75%. Clin. tests and studies have shown
     that, with use of the protonic mixture, about 30 to 40% more amino acids are
     utilized than when the protonic mixture is not used.
AN
     143:114482 CA
ΤI
    Protein formulation comprising enzymes and pH adjusters for improved
    bioavailability of amino acids
IN
    Ernest, Michael
    Doctor's Signature Sales and Marketing International Corp., USA
PA
SO
    U.S. Pat. Appl. Publ., 11 pp.
     CODEN: USXXCO
DT
     Patent
LΑ
    English
FAN.CNT 1
                       KIND DATE APPLICATION NO.
    PATENT NO.
                                                                 DATE
                        ----
                               -----
    US 2005152887
                        A1
                               20050714
                                           US 2004-757706
                                                                 20040114
PΙ
PRAI US 2004-757706
                               20040114
L3
    ANSWER 2 OF 13 CA COPYRIGHT 2005 ACS on STN
AB
    A complex enzyme composition that improves growth and feed digestion by animals
     comprises pancreatin 200, betaine-HCl 50, monobasic
     calcium phosphate 100, \alpha-amylase 150, \beta-amylase 100, lipase 50,
    pepsin 100, and cellulase 100 mg.
AN
     142:218044 CA
ΤI
    Complex enzyme composition containing pancreatin, betaine-
    HCl, and calcium phosphate as feed additive
IN
    Han, In Kyu
PΑ
    S. Korea
    Repub. Korean Kongkae Taeho Kongbo, No pp. given
SO
     CODEN: KRXXA7
DT
    Patent
    Korean
LΑ
FAN.CNT 1
                       KIND DATE
                                          APPLICATION NO. DATE
    PATENT NO.
    KR 2002041162
                        Α
                               20020601
                                           KR 2000-70943
                                                                  20001127
PΙ
PRAI KR 2000-70943
                               20001127
```

A composition containing betaine hydrochloride and pepsin as powders, AB pancreatin, α -amylase, β -amylase, lipase, cellulase, dibasic calcium phosphate as an enzyme complex, Lactobacillus acidophilus, Bifidobacteria longum, fructooligosaccharide or the like as a probiotics complex, a vitamin complex and a mineral complex is provided which inhibits proliferation of intestinal harmful microorganisms and enhances immunoactivity. The composition contains 78 mg betaine HCl 50 mg pepsin, 15 mg pancreatin, 11 mg α -amylase, 8 mg β -amylase, 4 mg lipase, 7.5 mg cellulase, 7.5 mg dibasic calcium phosphate, 1.25 billon Lactobacillus acidophilus, Lactobacillus bulgaricus, etc. 900 million Bifidobacteria longum and Bifidobacterium breve, 180 million Streptococcus thermophilus, 375 mg fructooligosaccharide, 450 IU vitamin A, 675 IU β -carotene, 56 mg vitamin C, 23 IU vitamin D, 23 IU vitamin E, 7.5 µg vitamin K, 3.8 mg thiamine, 3.4 mg riboflavin, 5.6 mg niacin, 3.4 mg pyridoxine, 11 µg cobalamin, 28 µg biotin, 8.4 mg pantothenic acid, 8.4 mg choline, 28.1 mg Ca, 18.8 mg P, 0.9 mg Fe, 12 μ g I, 17 mg Mg, 1.7 mg Zn, 11 μ g Se, 0.1 mg sulfuric acid, 0.6 mg Mn, 11 μ g chromium picolinate, 0.6 mg Mo, 0.6 mg K, 0.3 mg betaine, 0.2 mg B, 3.8 mg L-lysine 2.0 mg phenylalanine, 2.0 mg L-tyrosine, a trace amount of kelp powder and 4.7 mg polyunsatd. fatty acid.

AN 142:204717 CA

TI Nutrients containing betaine, enzymes, Lactobacillus, Bifidobacteria, fructooligosaccharides, vitamins and minerals

IN Han, In Kyu; Kim, Yu Yong

PA S. Korea

SO Repub. Korean Kongkae Taeho Kongbo, No pp. given

CODEN: KRXXA7

DT Patent

LA Korean

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
PI KR 2003025587	A	20030329	KR 2001-58713	20010921	
PRAI KR 2001-58713		20010921			

L3 ANSWER 4 OF 13 CA COPYRIGHT 2005 ACS on STN

AΒ A protonic formulation is provided comprising a protein mixture and a mix. of enzymes and pH adjusters selected for proper activation, pH adjustment, and attainment of pK for the amino acids and optimization of bioavailability of the amino acids. The optimum ratio of enzyme activator formulation to protein mixture is about 1:25, though 1 part enzyme activator formulation to 10-30 parts protein mixture will function suitably for the intended purpose. The enzyme activator formulation is optimally comprised of: betaine-HCl 4.0%, pepsin 1.5%, trypsin 0.4%, chymotrypsin 0.3%, protease 0.4%, bromelain 0.5%, papaya 0.6%, vitamin C 5.0%, lemon powder 0.6%, glutamic acid 0.2%, and glycine 86.4%. The protein sources and mixture that works best with the enzyme activator formulation includes: whey protein isolate 30.0%, instant whey concentrate 15.0%, soy protein isolate 25.0%, pea protein 5.0%, rice protein 5.0%, maltodextrin 15.7%, steviocide 0.3%, french vanilla flavor 1.75%, peach mango flavor 0.5%, xanthan gum 0.5%, lecithin 0.5%, and tricalcium phosphate 0.75%. Clin. tests have shown that, with use of the protonic mixture, about 30-40% more amino acids are utilized than when the protonic mixture is not used.

AN 138:169167 CA

TI Protein formulation with enzymes and pH adjusters for improved bioavailability of amino acids

IN Ernest, Michael

PA Life Force International, USA

SO PCT Int. Appl., 21 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN. CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003014304	A2	20030220	WO 2002-US24662	20020802

```
20030501
    WO 2003014304
                         Α3
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
             PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
             UA, UG, US, UZ, VN, YU, ZA, ZM, ZW
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
             FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF,
             CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
                                20010809
PRAI US 2001-311280P
                          Ρ
    ANSWER 5 OF 13 CA COPYRIGHT 2005 ACS on STN
T.3
    An herbal formulation useful as a food supplement for re-establishing
AΒ
     intestinal bacteria and rebuilding intestinal mucosa comprises 25-35%
    betaine HCl, 2-7% plant enzymes, 1-4% papain, 0.5-5%
    probiotic micro flora, 2-7% fructooligosaccharides, 5-15% L-glutamine,
     2-7% quercitin, 2-7% butyric acid, 5-15% borage seed, 5-15% flax seed,
     5-10% lecithin, and 5-15% of a mixture containing \gamma-oryzanol, bromelain,
     pepsin, and N-acetylglucosamine. The formulation may be mixed
     together, compressed and formed into a capsule for oral administration.
     137:129879 CA
AN
    Herbal formulation containing enzymes for rebuilding intestinal bacteria
ΤI
     Terry, Travis L.; Watson, Tommy Stanley; Watson, Brenda F.
IN
     Renew Life, Inc., USA
PΑ
SO
     U.S., 3 pp.
     CODEN: USXXAM
DT
     Patent
LΑ
     English
FAN. CNT 1
                                                                   DATE
                         KIND
                                DATE
                                            APPLICATION NO.
     PATENT NO.
                                                                    _____
     -----
     US 6426099
                          B1
                                20020730
                                            US 1998-204036
                                                                   19981201
PΙ
PRAI US 1997-67271P
                         Р
                                19971203
              THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 4
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 6 OF 13 CA COPYRIGHT 2005 ACS on STN
L3
     The therapeutic efficacy of dexamethasone and a natural pig surfactant
AB
     preparation was investigated in a rabbit aspiration model. Lung injury was
     induced by intratracheal administration of 2 mL of a betaine-
     HCl-pepsin mixture/kg. Dexamethasone was given i.v. in
     two doses (D1 = 7.5 \text{ mg/kg}; D2 = 3.75 \text{ mg/kg}; D2 6 h post D1). In different
     groups D1 was injected at different times before and after aspiration.
     Natural surfactant was administered 24 h post lung injury in a single dose
     of 12 mg phospholipids/kg. The therapeutic potential was evaluated by
     measuring static lung compliance and the difference in a lung volume between
     0 and 20 mm Hg airway pressure. No therapeutic effect of dexamethasone
     was seen at any time of application. In contrast, the intratracheal
     administration of natural surfactant 24 h post injury completely reversed
     the deterioration of lung mech. properties.
     119:86523 CA
AN
     Experimental aspiration trauma: Comparison of steroid treatment versus
TΤ
     exogenous natural surfactant
ΑU
     Strohmaier, W.; Schlag, G.
     Ludwig Boltzmann Inst. Exp. Clin. Traumatol., Vienna, Austria
CS
     Experimental Lung Research (1993), 19(3), 397-405
SO
```

L3 ANSWER 7 OF 13 CA COPYRIGHT 2005 ACS on STN

CODEN: EXLRDA; ISSN: 0190-2148

DT

LΑ

Journal

English

AB Pharmaceuticals for treatment of digestive tract disorders in domestic animals contain betaine-HCl 10-30, antacid carbohydrate digestive enzymes 1-10, antacid cellulose-degrading enzymes 1-10, antacid protein digestive enzymes 20-40, and saccharification bacteria spore powder 10-30% by weight Thus, a pharmaceutical was prepared by

mixing betaine-HCl 200, carbohydrate digestive enzyme 50, a cellulose degrading enzyme 50, sugar-containing pepsin 300, saccharification bacterial spore powder 200, lactose 100 parts by weight, and potato starch q.s.

AN 112:62655 CA

TI Pharmaceuticals for treatment of digestive tract disorders of domestic animals

IN Masuda, Takashi

PA Toa Yakuhin Kogyo K: K., Japan

SO Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	JP 01132533	A2	19890525	JP 1987-289628	19871118
PRAT	TP 1987-289628		19871118		

L3 ANSWER 8 OF 13 CA COPYRIGHT 2005 ACS on STN

- The characteristic enzymic activity, using the gastrointestinal model, of the acid-resistant digestive enzyme which had been added to the preparation containing betaine-HCl was determined and compared with those of folk digestive enzyme prepns. The deviation of the wts. of those prepns. was also investigated. The present preparation shows it digestive activity in the acid range and act in the stomach. The digestive activity using the shaking methods and the separated methods, i.e. amylase, protease and lipase did not fulfill the standard activity of the digestive enzyme prepns. The digestion using the gastrointestinal model was about the same as that of the digestive enzyme prepns. fulfilling th criteria and, especially in the gastric model, it was same or more than that. The prepns. in which more than 2 types of granules, had been mixed fulfilled the requirements of the weight variation in the Japanese Pharmacopoeis, but the mixture ratio of those was variable.
- AN 109:176308 CA
- TI The characteristic digestive activity of the preparation containing betaine hydrochloride
- AU Murakami, Tadayasu; Kawashima, Mikio; Sasaki, Masanori; Kobayashi, Shinichi; Yamada, Fusayo; Asahina, Kikuo
- CS Res. Lab., Toa Pharm. Co., Ltd., Tatebayashi, 374, Japan
- SO Yakuri to Chiryo (1973-2000) (1988), 16(2), 771-8 CODEN: YACHDS; ISSN: 0386-3603
- DT Journal
- LA Japanese
- L3 ANSWER 9 OF 13 CA COPYRIGHT 2005 ACS on STN
- Oval compns. for decreasing the symptoms of digestive dysfunction contain a pancreatic enzyme extract, a protelytic enzyme from plants, a choleretic agent, a HCl source and pepsin [9001-75-6]. Bromelain [9001-00-7] and pancreatin [8049-47-6] were adsorbed onto digestible sucrose-starch beads which were coated with while lac glaze. These beads were then coated with stearic acid and carnauba wax. A mixture of ox bile extract, pepsin, betaine-HCl [590-46-5] and guar gum was blended with H2O to give a dough which was screened, dried, and the resultant granules ground and dry-screened to the mesh. The granules were mixed with the coated beads and blended with hydrogenated vegetable oil, microcryst. cellulose, and Mg stearate. The mixture was punched into tablets and coated with a zein solution
- AN 101:28325 CA
- TI Enzyme-containing digestive aid compositions
- IN Bilton, Gerald L.
- PA USA
- SO U.S., 4 pp.
- CODEN: USXXAM
- DT Patent
- LA English
- FAN.CNT 1

```
_____
                        ____
                               _____
                                           ------
                             19840508 US 1983-462995
                                                                  19830201
    US 4447412 ·
                       Α
                                                                 19840203
                        A1 19850815 WO 1984-US159
    WO 8503438
        W: JP
        RW: AT, BE, CH, DE, FR, GB, LU, NL, SE
                        A1 19860226 EP 1984-901130
                                                                 19840203
    EP 172166
        R: AT, BE, CH, DE, FR, GB, LI, LU, NL, SE
    CA 1213543 A1 19861104 CA 1984-447208
                                                                19840210
                               19830201
PRAI US 1983-462995
                        Α
                               19840203
    WO 1984-US159
    ANSWER 10 OF 13 CA COPYRIGHT 2005 ACS on STN
L3
    A review with refs. of betaine-HCl [590-46-5],
AB
    glutamic acid-HCl [138-15-8], diluted HCl, and pepsin
     [9001-75-6] as ingredients in over-the-counter (OTC) drug products for use
    as stomach acidifiers. Based upon the lack of adequate data to establish
     the effectiveness of these or any other ingredients of stomach acidifiers
    used in treating achlorhydria and hypochlorhydria, and because such
     conditions are asymptomatic and not ameniable to self-diagnosis, any OTC
    drug product containing ingredients offered for use as stomach acidifiers
    cannot be considered generally recognized as safe and effective.
AΝ
    Stomach acidifier drug products for over-the-counter human use; proposed
ТT
    rulemaking
    Food and Drug Administration, Rockville, MD, 20857, USA
CS
     Federal Register (1979), 44(204), 60316-20, 19 Oct 1979
SO
     CODEN: FEREAC; ISSN: 0097-6326
DT
     Journal; General Review
LΑ
    English
    ANSWER 11 OF 13 CA COPYRIGHT 2005 ACS on STN
L3
    A combination of 455 mg. betaine-HCl and 60 mg.
AB
    pepsin (1:10,000 U.S.P. unit), having the mixed powder particles
    coated with 141 mg. methylcellulose, is placed in capsules. The mixture is useful as a gradual producer of HCl in patients with achlorhydria or
     hypochlorhydria. Glutamic acid-HCl can replace the betaine-
     HCl.
AN
     51:83316 CA
OREF 51:15073b-c
     Preparation containing betaine hydrochloride for treatment of achlorhydria
     and hypochlorhydria
     Sahyun, Melville
TN
DT
     Patent
LA
    Unavailable
     PATENT NO. KIND DATE APPLICATION NO. DATE
FAN.CNT 1
    PATENT NO.
                               19570709 US
     US 2798837
PI
L3
    ANSWER 12 OF 13 CA COPYRIGHT 2005 ACS on STN
     Betaine unites with HCl loosely to form a compound which readily breaks up
AB
     into its components in aqueous solution Because of this property the substance
     forms a convenient medium for the administration of HCl. Betaine
     -HCl contains 23.8% of HCl. Acidol is a proprietary name for
     the substance and its mixts. with pepsin are called "acidol-
     pepsin." Betaine-HCl is a white, crystalline,
     odorless substance of an acid reaction and taste. About 10 yrs. ago the
     acid contents of acidol and of acidol-pepsin were determined The
     amts. found were substantially as claimed. The acid was determined by titration
     with N KOH, using phenolphthalein as indicator, and by precipitation with AgNO3
     and weighing as AqCl. Recently new specimens of each product were examined
     The old products were reexamd. and the results compared. The acidity and
     the proteolytic activity of the new specimens were essentially as claimed.
     The acidity of the old specimens had not changed much in 10 yrs., but the
     proteolytic activity had disappeared.
     14:18796 CA
AN
```

OREF 14:3501b-d

Acidol and acidol-pepsin

```
AII
     Anon.
SO
     Rep. Lab. Am. Med. Assoc. (1919), 12, 91-3
DT
     Journal
LA Unavailable
L3
     ANSWER 13 OF 13 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on
AB
     An herbal formulation comprises betaine HCl, plant
     enzymes, papain, probiotic micro flora, fruitooligosaccharides,
     1-glutamine, quercitin, butyric acid, borage seed, flax seed, lecithin,
     gamma oryzanol, bromelain, pepsin, and N-acetylglucosamine.
     2002:477375 BIOSIS
AN
DN
     PREV200200477375
     Herbal formulation for rebuilding intestinal bacteria.
ΤI
ΙΙΔ
     Terry, Travis L. [Inventor, Reprint author]; Watson, Tommy Stanley
     [Inventor]; Watson, Brenda F. [Inventor]
CS
     Clearwater, FL, USA
     ASSIGNEE: Renew Life, Inc., Clearwater, FL, USA
     US 6426099 20020730
PΙ
     Official Gazette of the United States Patent and Trademark Office Patents,
SO
     (July 30, 2002) Vol. 1260, No. 5. http://www.uspto.gov/web/menu/patdata.ht
     ml. e-file.
     CODEN: OGUPE7. ISSN: 0098-1133.
DT
     Patent
LA
     English
ED
     Entered STN: 11 Sep 2002
     Last Updated on STN: 11 Sep 2002
=> s whey protein
          9182 WHEY PROTEIN
L4
=> s soy protein
L5
          8269 SOY PROTEIN
\Rightarrow s 14 and 15
           276 L4 AND L5
L6
=> s 14 (p) 15
L7
           249 L4 (P) L5
=> d his
     (FILE 'HOME' ENTERED AT 18:34:18 ON 14 OCT 2005)
     FILE 'CA, BIOSIS, MEDLINE' ENTERED AT 18:36:52 ON 14 OCT 2005
            242 S (BETAINE HCL) OR (BETAINE HYDROCHLORIC ACID)
L1
L2
          41097 S PEPSIN
L3
             13 S L1 AND L2
           9182 S WHEY PROTEIN
L4
L5
           8269 S SOY PROTEIN
            276 S L4 AND L5
1.6
L7
            249 S L4 (P) L5
=> s 11 and 12 and 14 and 15
             2 L1 AND L2 AND L4 AND L5
=> d 1-2 ab, bib
     ANSWER 1 OF 2 CA COPYRIGHT 2005 ACS on STN
L8
AΒ
     The present invention provides a stabilized protonic formulation comprised
     primarily of proteins, enzymes and pH adjusters, all in specific ratios to
     one another, a liquid medium which, when combined to the protonic
     formulation, initiates activation of the amino acids within the protonic
     formulation, and a stabilizing component which stabilizes the amino acids
     during the process of their activation. The optimum ratio of enzyme
     activator formulation to protein mixture is about 1:25, though 1 part enzyme
```

activator formulation to 10 to 30 parts protein mixture will function

suitably for the intended purpose. The enzyme activator formulation is optimally comprised of: betaine-HCl 4.0%, pepsin 1.5%, trypsin 0.4%, chymotrypsin 0.3%, protease 0.4%, bromelain 0.5%, papaya 0.6%, vitamin C 5.0%, lemon powder 0.6%, glutamic acid 0.2%, and glycine 86.4%. The protein sources and mixture that works best with the enzyme activator formulation includes: whey protein isolate 30.0%, instant whey concentrate 15.0%, soy protein isolate 25.0%, pea protein 5.0%, rice protein 5.0%, maltodextrin 15.7%, steviocide 0.3%, french vanilla flavor 1.75%, peach mango flavor 0.5%, xanthan gum 0.5%, lecithin 0.5%, and tricalcium phosphate 0.75%. Clin. tests and studies have shown that, with use of the protonic mixture, about 30 to 40% more amino acids are utilized than when the protonic mixture is not used. 143:114482 CA Protein formulation comprising enzymes and pH adjusters for improved bioavailability of amino acids Ernest, Michael Doctor's Signature Sales and Marketing International Corp., USA U.S. Pat. Appl. Publ., 11 pp. CODEN: USXXCO Patent English FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE ----US 2005152887 US 2004-757706 A1 20050714 20040114 PRAI US 2004-757706 20040114 ANSWER 2 OF 2 CA COPYRIGHT 2005 ACS on STN A protonic formulation is provided comprising a protein mixture and a mix. of enzymes and pH adjusters selected for proper activation, pH adjustment, and attainment of pK for the amino acids and optimization of bioavailability of the amino acids. The optimum ratio of enzyme activator. formulation to protein mixture is about 1:25, though 1 part enzyme activator formulation to 10-30 parts protein mixture will function suitably for the intended purpose. The enzyme activator formulation is optimally comprised of: betaine-HCl 4.0%, pepsin 1.5%, trypsin 0.4%, chymotrypsin 0.3%, protease 0.4%, bromelain 0.5%, papaya 0.6%, vitamin C 5.0%, lemon powder 0.6%, glutamic acid 0.2%, and glycine 86.4%. The protein sources and mixture that works best with the enzyme activator formulation includes: whey protein isolate 30.0%, instant whey concentrate 15.0%, soy protein isolate 25.0%, pea protein 5.0%, rice protein 5.0%, maltodextrin 15.7%, steviocide 0.3%, french vanilla flavor 1.75%, peach mango flavor 0.5%, xanthan gum 0.5%, lecithin 0.5%, and tricalcium phosphate 0.75%. Clin. tests have shown that, with use of the protonic mixture, about 30-40% more amino acids are utilized than when the protonic mixture is not used. 138:169167 CA Protein formulation with enzymes and pH adjusters for improved bioavailability of amino acids Ernest, Michael Life Force International, USA PCT Int. Appl., 21 pp. CODEN: PIXXD2 Patent English FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE -------------------WO 2003014304 A2
WO 2003014304 A3 20030220 WO 2002-US24662 20020802 WO 2003014304 A3 20030501 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,

AN

ΤI

IN

PA SO

DT

LΑ

PΙ

L8

AΒ

AN

TΤ

IN

PA

SO

DT

LΑ

PΙ

```
KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI US 2001-311280P P 20010809
```

=> d his

(FILE 'HOME' ENTERED AT 18:34:18 ON 14 OCT 2005)

	FILE 'CA, BIOSIS, MEDLINE' E	ENTERED AT 18:36:52 ON 14 OCT 2005
L1	242 S (BETAINE HCL) O	R (BETAINE HYDROCHLORIC ACID)
L2	41097 S PEPSIN	
L3	13 S L1 AND L2	
L4	9182 S WHEY PROTEIN	
L5	8269 S SOY PROTEIN	
L6	276 S L4 AND L5	
L7	249 S L4 (P) L5	
L8	2 S L1 AND L2 AND L	4 AND L5

(FILE 'HOME' ENTERED AT 18:34:18 ON 14 OCT 2005)

	FILE 'CA, BIOSIS, MEDLINE	' ENTERED AT 18:36:52 ON 14 OCT 2005
L1	242 S (BETAINE HCL	OR (BETAINE HYDROCHLORIC ACID)
L2	41097 S PEPSIN	
L3	13 S L1 AND L2	
L4	9182 S WHEY PROTEIN	
L5	8269 S SOY PROTEIN	
L6	276 S L4 AND L5	
L7	249 S L4 (P) L5	
L8	2 S L1 AND L2 AN	D L4 AND L5
L9	3128 S NERVOUS SYST	EM DISORDERS
L10	0 S L7 AND L9	
L11	0 S AUTISM AND L	7

Freeform Search

	US Pat US OC EPO A JPO AI Derwei	tents Full-Te R Full-Text bstracts Dat bstracts Dat nt World Pat	labase abase		se				
Term:	17 an	d 120		•••••	,			,	
Display:			its in <u>Displa</u> y				g with N	lumber	20
Generate:	: C Hit	: List © B	Search	Side by		Image			
	•••••		······						·····

DATE: Friday, October 14, 2005 Printable Copy Create Case

Set Name side by side	Query	Hit Count	Set Name result set
DB=PGPB	,USPT,USOC,EPAB,JPAB,DWPI,TDBD; PLUR=Y	ES; OP=ADJ	
<u>L21</u>	17 and 120	45	<u>L21</u>
<u>L20</u>	probiotic	1982	<u>L20</u>
<u>L19</u>	17 and 115	0	<u>L19</u>
<u>L18</u>	17 and 117	0	<u>L18</u>
<u>L17</u>	autism	2661	<u>L17</u>
<u>L16</u>	L15 and 18	0	<u>L16</u>
<u>L15</u>	nervous system disorder	7489	<u>L15</u>
<u>L14</u>	18 and autism	0	<u>L14</u>
<u>L13</u>	11 and 12 and 15 and 16	1	<u>L13</u>
<u>L12</u>	110 and 19 and 11 and 12	1	<u>L12</u>
<u>L11</u>	19 same 110	126	<u>L11</u>
<u>L10</u>	soy protein isolate	1489	<u>L10</u>
<u>L9</u>	whey protein isolate	538	<u>L9</u>
<u>L8</u>	15 same 16	949	<u>L8</u>
<u>L7</u>	15 and 16	1267	<u>L7</u>
<u>L6</u>	soy protein	6505	<u>L6</u>

<u>L5</u>	whey protein	5285	<u>L5</u>
<u>L4</u>	whey protein isolate	538	<u>L4</u>
<u>L3</u>	11 and 12	13	<u>L3</u>
<u>L2</u>	pepsin	21063	<u>L2</u>
<u>L1</u>	betaine hcl or betaine hydrochloric acid	78	<u>L1</u>

END OF SEARCH HISTORY